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- (5) With the oxides of nitrogen analyzer in the NO mode, record the concentration of NO indicated by the analyzer.
- (6) Turn on the NO_X generator O_2 (or air) supply and adjust the O_2 (or air) flow rate so that the NO indicated by the analyzer is about 10 percent less than indicated in paragraph (b)(5) of this section. Record the concentration of NO in this $NO+O_2$ mixture as value "c."
- (7) Switch the NO_X generator to the generation mode and adjust the generation rate so that the NO measured on the analyzer is 20 percent of that measured in paragraph (b)(5) of this section. There must be at least 10 percent unreacted NO at this point. Record the concentration of residual NO as value "d"
- (8) Switch the oxides of nitrogen analyzer to the NO_X mode and measure total NO_X . Record this value as "a."
- (9) Switch off the NO_X generator but maintain gas flow through the system. The oxides of nitrogen analyzer will indicate the NO_X in the $NO+O_2$ mixture. Record this value as "b".
- (10) Turn off the NO_X generator O_2 (or air) supply. The analyzer will now indicate the NO_X in the original NO-in-N2 mixture. This value should be no more than five percent above the value indicated in paragraph (b)(4) of this section.
- (11) Calculate the efficiency of the $NO_{\rm X}$ converter by substituting the concentrations obtained into the following equation:

percent efficiency =
$$\left(1 + \frac{a - b}{c - d}\right) \times 100$$

Where:

- a = concentration obtained in paragraph
 (b)(8),
- b = concentration obtained in paragraph (b)(9).
- c = concentration obtained in paragraph
 (b)(6),
- d = concentration obtained in paragraph
 (b)(7).

If converter efficiency is less than 90 percent, corrective action will be required.

(c) Initial and periodic calibration. Prior to its initial use and monthly thereafter, or within one month prior

to the certification test, calibrate the chemiluminescent oxides of nitrogen analyzer on all normally used instrument ranges. Use the same flow rate as when analyzing samples. Proceed as follows:

- (1) Adjust analyzer to optimize performance.
- (2) Zero the oxides of nitrogen analyzer with purified synthetic air or zero-grade nitrogen.
- (3) Calibrate on each normally used operating range with NO-in-N₂ calibration gases having nominal concentrations between 10 and 90 percent of that range. A minimum of six evenly spaced points covering at least 80 percent of the 10 to 90 range (64 percent) is required (see following table).

Example calibration points (%)	Acceptable for calibration?
20, 30, 40, 50, 60, 70	No, range covered is 50 percent, not 64 Yes. Yes. No, though equally spaced and entire range covered, a minimum of six points are needed.

Additional calibration points may be generated. For each range calibrated, if the deviation from a least-squares best-fit straight line is two percent or less of the value at each data point, calculate concentration values by use of a single calibration factor for that range. If the deviation exceeds two percent at any point, use the best-fit non-linear equation which represents the data to within two percent of each test point to determine concentration.

(d) The initial and periodic interference, system check, and calibration test procedures specified in 40 CFR part 1065, subpart D, may be used in lieu of the procedures specified in this section.

[60 FR 34598, July 3, 1995, as amended at 70 FR 40449, July 13, 2005]

$\S 90.319$ NO $_X$ converter check.

- (a) The efficiency of the converter used for the conversion of NO_2 to NO is tested as given in paragraphs (a)(1) through (a)(8) of this section.
- (1) Using the test setup as shown in Figure 1 in Appendix B of this subpart (see also §90.318 of this chapter) and the procedure described in paragraphs (a)(2) through (a)(8) of this section, test

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the efficiency of converters by means of an ozonator.

- (2) Calibrate the HCLD or CLD in the most common operating range following the manufacturer's specifications using zero and span gas (the NO content of which must amount to about 80 percent of the operating range and the $\rm NO_2$ concentration of the gas mixture less than five percent of the NO concentration). The $\rm NO_X$ analyzer must be in the NO mode so that the span gas does not pass through the converter. Record the indicated concentration.
- (3) Calculate the efficiency of the NO_X converter as described in §90.318(b).
- (4) Via a T-fitting, add oxygen continuously to the gas flow until the concentration indicated is about 20 percent less than the indicated calibration concentration given in paragraph (a)(2) of this section. Record the indicated concentration "c." The ozonator is kept deactivated throughout the process.
- (5) Activate the ozonator to generate enough ozone to bring the NO concentration down to about 20 percent (minimum 10 percent) of the calibration concentration given in paragraph (a)(2) of this section. Record the indicated concentration "d."

Note: If, with the analyzer in the most common range, the $NO_{\rm X}$ converter can not give a reduction from 80 percent to 20 percent, then use the highest range which will give the reduction.

- (6) Switch the NO analyzer to the NO_X mode which means that the gas mixture (consisting of NO, NO_2 , O_2 and N_2) now passes through the converter. Record the indicated concentration "a."
- (7) Deactivate the ozonator. The mixture of gases described in paragraph (a)(6) of this section passes through the converter into the detector. Record the indicated concentration "b."
- (8) Switched to NO mode with the ozonator deactivated, the flow of oxygen or purified synthetic air is also shut off. The NO_X reading of the analyzer may not deviate by more than \pm five percent of the theoretical value of the figure given in paragraph (a)(2) of this section.

- (b) The efficiency of the converter must be tested prior to each calibration of the NO_X analyzer.
- (c) The efficiency of the converter may not be less than 90 percent.

§ 90.320 Carbon dioxide analyzer calibration.

- (a) Prior to its initial use and monthly thereafter, or within one month prior to the certification test, calibrate the NDIR carbon dioxide analyzer as follows:
- (1) Follow good engineering practices for instrument start-up and operation. Adjust the analyzer to optimize performance.
- (2) Zero the carbon dioxide analyzer with either purified synthetic air or zero-grade nitrogen.
- (3) Calibrate on each normally used operating range with carbon dioxide-in- N_2 calibration or span gases having nominal concentrations between 10 and 90 percent of that range. A minimum of six evenly spaced points covering at least 80 percent of the 10 to 90 range (64 percent) is required (see following table).

Example calibration points (%)	Acceptable for Calibration?
20, 30, 40, 50, 60, 70	No, range covered is 50 percent, not 64.
20, 30, 40, 50, 60, 70, 80, 90	Yes.
10, 25, 40, 55, 70, 85	Yes.
10, 30, 50, 70, 90	No, though equally spaced and entire range covered, a minimum of six points are needed.

Additional calibration points may be generated. For each range calibrated, if the deviation from a least-squares best-fit straight line is two percent or less of the value at each data point, calculate concentration values by use of a single calibration factor for that range. If the deviation exceeds two percent at any point, use the best-fit non-linear equation which represents the data to within two percent of each test point to determine concentration.

(b) The initial and periodic interference, system check, and calibration test procedures specified in 40 CFR part 1065, subparts C and D, may be used in lieu of the procedures in this section.

[60 FR 34598, July 3, 1995, as amended at 70 FR 40449, July 13, 2005]